

EUPATI PATIENT EXPERT TRAININGS

The Patient Expert Programmes aim to educate patients on the medicines R&D process and their role. National programs offer territorial and national perspectives.



The Netherlands

EUPATINL Fellow



Open to everyone interested in Medicines R&D.
Motivation letter to register

Training in English

Free of charge

250 hours of online learning + 8 training days (online)



Sweden



The web-based training features four courses, each taking **2-3 hours with a total of 12-hour** training.
.Provided in Swedish. Open to everyone



Ireland

IPPOSI graduate



Open to Patients, P. Representatives and Careres.
Motivation letter and Online form to register

Training in English

Free of charge

160 h of online learning + 4x1/2 in person days + 6 x1/2 virtual days



Czech Republic



Course of 6 one-day training events, each lasting 8 hours, for a **total of 48 hours**. Provided in Czech.
Open to everyone .



Switzerland



Course of 6 online self-study modules + 1 day in person per module. Provided in German. Open to everyone **Basic: 8-16 h Additional modules: 16-32 h**



Spain



Course **46 hours** (15 h online and 12h face-to-face).
Provided in Spanish. Open to patient organization representatives



Italy

Eupati Patient Expert



Open to Patients, P. Representatives and Careres.
Motivation letter, CV and online form to register

Training in Italian

Free of charge

160 h of online learning + 6 in person training days



Cyprus



Course of 7 in person sessions lasting 3 hours each, **totaling 21 hours** over 7 weeks. Provided in Greek.
Open to Patients, careers, and their advocates



PATIENT EXPERT TRAINING PATHWAYS - LONG TRAININGS



	PATIENT EXPERT TRAINING PROGRAMME EUPATI Fellow	DUTCH NATIONAL PLATFORM PATHWAY EUPATINL Fellow	IRISH NATIONAL PLATFORM PATHWAY IPPOSI graduate	ITALIAN NATIONAL PLATFORM PATHWAY Eupati Patient Expert
	Open to all interested in Medicines R&D and patient engagement, registration on the EUPATI website is on a first-come, first-served basis with registration open from June to 1st November. 130 participants per cohort.	Open to patients, representatives, or carers interested in Medicines R&D. Registration requires submitting a motivation letter and possibly a letter from a patient organization. Selection is done by a working group. 18 participants per cohort.	Open to patients, careers, and patient advocates. Register by submitting an online application and a motivation letter. Selection based on application score by a committee. 20-25 participants per cohort.	Open to patients, patient representatives, caregivers, and some other stakeholders. Applicants submit an online application with a CV and motivation letter for evaluation by a selection committee based on a score. 40 participants per cohort
	12-14 months training <ul style="list-style-type: none"> • 300 hours of online learning • Two 4-day-training events One online one in person 	14 months training <ul style="list-style-type: none"> • 250 hours of online learning • 8 training days (at least a training day per module) 	10 months training <ul style="list-style-type: none"> • 160 hours of online learning • 4 half days in-person + 6 half days virtual 	10 months training <ul style="list-style-type: none"> • 160 hours of online learning • 6 training days
	Training provided in English	Training provided in English	Training provided in English	Training provided in Italian
	<ul style="list-style-type: none"> • Getting started • Introduction to Medicines R&D • Non-Clinical Development • Clinical Development • Regulatory Affairs • Health Technology Assessment (HTA) 	<ul style="list-style-type: none"> • Process of Medicines Discovery and development • Non-Clinical Development • Clinical Development • Registration, safety and pharmacovigilance • Health Technology Assessment (HTA) and reimbursement 	<ul style="list-style-type: none"> • Clinical Trials • Medicine & Medical Device Regulation • Health Technology Assessment 	<ul style="list-style-type: none"> • Drug discovery and development planning • Pre-clinical phase and drug development • Exploratory and confirmatory clinical development • Clinical studies • Regulatory Affairs, Safety of Medicines, PV and Pharmacoepidemiology • Principles and practice of HTA
	208€ + Travel expenses for in person event. Limited scholarships available	Free of charge	Free of charge	Free of charge

PATIENT EXPERT TRAINING PATHWAYS - SHORT COURSES



	 SWISS NATIONAL PLATFORM TRAINING EUPATI CH Patientenexperte	CZECH NATIONAL PLATFORM TRAINING Czech ENP/APO participant	SWEDISH NATIONAL PLATFORM TRAINING EUPATI Sweden expert patient	CYPRUS NATIONAL PLATFORM TRAINING EUPATI Patient Expert	SPANISH NATIONAL PLATFORM TRAINING
	Open to all interested in Medicine R&D via online registration form with 20-25 participants per cohort.	Open to all interested in Medicine R&D via online registration form with 15 participants per cohort.	Anyone with an interest in medical research. Registration by creating an account on their learning platform. No limit of participants	Open to Patients, careers, and their advocates via online registration with 27 participants per cohort.	Open to patient organization representatives seeking to learn about medicine R&D for educating their trainees via online registration form with 20 participants per cohort.
	The course consists of 6 modules with online self-study materials and one-day in person training with experts for each module. Basic course requires 8-16 hours over 2 days , additional modules need 16-32 hours over 4 days . Training spans 8 months.	The training aims to localize EUPATI core modules into Czech with expert guarantors overseeing the creation of aligned lectures and translation of key terms from the EUPATI Toolbox. It consists of 6 one-day training events (8 hours each), totaling 48 hours , spread over 6 months.	The web-based training includes four courses tailored to Swedish conditions with expert lectures, readings, and assignments. Each course takes 2-3 hours, totaling around 12-15 hours for the entire training , which can be completed at the learner's own pace.	The training consists of 7 in person sessions lasting 3 hours each, totaling 21 hours over 7 weeks . Including presentations and a short online test after completion.	Course consisting of an online part of 46 hours including 15 hours of virtual classes (via zoom) and a face-to-face part of 12 hours .
	Training provided in German	Training provided in Czech	Training provided in Swedish	Training provided in English and Greek	Training provided in Spanish
	The training covers human research fundamentals in Switzerland: ethical and legal aspects, research priorities, study methodology, implementation of ethical and legal requirements, Information and communication, and transferring new findings into practice	The training covers EUPATI courses adapted to the Czech including Introduction to medicines R&D, non-clinical development, Clinical Development, regulatory affairs, and Health Technology Assessment (HTA).	The programme is structured in four parts: This includes the journey of medicines from idea to patient, human rights, research and development, evaluation for approval, and marketing.	Training in personal development, Introduction to research, patient advocacy, drug development and Licensing, know-how/expertise, national strategies and HTA.	Drug discovery and development planning, pre-clinical phase and drug development, exploratory and confirmatory clinical development, clinical studies, regulatory affairs, Safety of medicines, PV and Pharmacoeconomics and principles and practice of HTA
	Partially covered (Total cost of CHF 3,300) participant covers CHF 1000 for all the modules	Free of charge	Free of charge	Free of charge	Free of charge